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 $\langle f|T|cell$ peptide epitopes are not associated with tronchial mucosal infiltration of eosinophils of TH 1 type cells or with elevated concentrations of histamine or ercosanoids in bronchoalveolar fluid.

Haselden B M; Larche M; Meng C; Shirley

F; Dworski R; Kaplan A P: Bates C; Robinson D B; Ying S;

AUTHOR -

Kay A B Lepartment of Allergy and Clinical Immunology, National CORPORATE SOURCE:

Heart and Lung Institute Imperial College School of Medicine, London, United Kingdom. DM 15431 NIGMS:

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Last Updated on STN: 20011015
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BACKGROUND: Isolated late asthmatic reactions can be provoked by intradermal challenge of allergen derived T cell peptide epitopes.
OBJECTIVE: The propose of this study was to determine whether the isolated LAR is associated with the local accumilation of inflammatory cells, the expression of T-N-2 cytokines, and the production of pharmacologic mediators. METHOUS: A randomized, placebo controlled, crossover study design was used. The investigation involved bronchial and skin biopsies and bronchoalveolar lavage (BAL) fluids from 8 cat-allergic subjects who developed significant late asthmatic reactions 6 hours after intradermal injection of Fel d 1 chain 1 derived peptides (FCIPS: RESULTS: remunostaining of bronchial biopsy specimens showed no changes in the injection of Fel d 1 chain 1 derived peptides (FC1Ps). RESULTS:
Immunostaining of bronchial biopsy specimens showed no changes in the
numbers of eosinaphils, neutrophils basophils, mast cells, CD3.*, CD4.*.

or CD8.(*) T cells, CD25.(*) cells or macrophages, or cells mRNA.* for
12.4, IL.5, or IL.13 when the FC1P day was compared with the
diluent control day. There were also no significant differences in
eosinophil numbers, either in BAL fluids or in peripheral blood after
FC1P challenge. Furthermore, there were no significant alterations
in the concentrations of histamine, histamine releasing factors, or
erosanoids (LTC141/D(4)/E14), PCD.2), PCB(2), PCB(2), PGF(2alpha) in BAL
fluids. FC1Ps induced a significant (P <.05) elevation in CD8.*) cells in
the skin and an unexpected decrease in IL.5 in BAL fluids (F =.043).
CONCLUSION Part of the asthma process might involve T cell dependent
alreay narrowing with no requirement for IgE, mast cells, or infiltrating
inflammatory cells. inflammatory cells

L4 ANSWER 2 OF 2 ACCESSION NUMBER:

MEDLINE

DOCUMENT NUMBER TITLE:

199307274 MEDLINE 29307274 FubMed ID: 10377184 Immunoglobulin E-independent major histocompatibility complex restricted Ticell peptide epitope induced late asthmatic reactions

AUTHOR CORPORATE SOURCE:

Ascimatic reactions

Asselden B M; Kay A B; Larche M

Fepartment of Allergy and Clinical Immunology, National

Heart and Lung Institute, Imperial College School of

Medicine, London SNS 6LY, United Kingdom.

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ENTRY LATE:

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Entered Meddine: 19990726
Distradermal administration of short overlapping peptides derived from chain 1 of the dat allerge. Fel d 1 *FCIP* that did not cross link 1gE, elicited isolated late asthmatic reactions with no visible early or late chaneous response in 9,40 cat allergic asthmatics. Four of the nine were human histocompatibility leukocyte antigen DR13 positive, as compared with only 1/31 norreactors. The other five reactors expressed either DR1 or DR1. To confirm major histocompatibility complex restriction, fibroblast cell lines transfected with HLA DR molecules were used to present PCIPs to dat allergen specific T cell lines derived from subjects before peptide injection. FCIPs peptide 28 44 of Fel d 1 chain 1 was recognized in the context of DR13 alleles DRB1*1301, 1302 and induced specific T cell proliferation and IL 5 production. T cells from a DR1 *, responder proliferated and produced IL 5 in the presence of FCIP3 and DR1 DRB1*0101* fibroblast cell lines, whereas T cells from a DR1 *, responder proliferated and produced IL 5 in the presence of FCIP3 and DR1 DRB1*0101* fibroblast cell lines, whereas T cells from a DR1 *, subject recognized FCIP2 peptide 22 3" when presented by DRB1*0405. We conclude that short allergen derived peptides can directly initiate a major histocompatibility complex restricted. T cell dependent late authmatic reaction, without the requirement for an early 19F/mast cell dependent :esponse, in sensitized asthmatic subjects.

cell dependent response, in sensitized asthmatic subjects.

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DOCUMENT NUMBER: TITLE:

PREV199900134427

AUTHOR:S:: CORPORATE SOURCE: SOURCE:

PREV199900134427
Peptide induced late
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MHC restricted T cell activation in vivo.
Larche, M.; Haselden, B. M.; Kay, A. B.
Natl. Heart Lung Inst., Imperial Coll. Sch. Med., London UK
Journal of Allergy and Clinical Immunology, Jan., 1999
Vol. 103, No. 1 PART 2, pp. S204.
Meeting Info.: 55th Annual Meeting of the American Academy
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